Listing of Claims:

1-27. (Canceled)

28. (previously presented) A composition for delivering an agent to cells, the composition comprising the agent and a delivery enhancing compound of Formula I:

$$X_1$$
— C — N — $(CH_2)_m$ — N — $(CH_2)_n$ — N — R
 C = O
 I
 X_2

wherein:

m and n are the same or different and each is an integer from 2-8; R forms a cationic group with the nitrogen to which it is bound, or

 X_1 is selected from the group consisting of

 X_2 , and X_3 are each independently selected from the group consisting of a saccharide group,

wherein at least one of X₂ and X₃ is a saccharide group when R is

——C——X₃, and wherein said agent is a member selected from the group consisting of a therapeutic protein, a therapeutic genes, a vector

and an antisense nucleic acid.

- 29. (previously presented) The composition according to claim 28, wherein the saccharide group has between one to eight monosaccharide groups.
- 30. (original) The composition according to claim 29, wherein the saccharide group is selected from the group consisting of pentose monosaccharide groups, hexose monosaccharide groups, pentose-pentose disaccharide groups, hexose-hexose disaccharide groups, and hexose-pentose disaccharide groups.
- 31. (original) The composition according to claim 28, wherein the saccharide group is a trisaccharide.
- 32. (original) The composition according to claim 28, wherein the concentration of the delivery enhancing compound is about 0.002 to about 2 mg/ml.

- 33. (original) The composition according to claim 32, wherein the concentration of the delivery enhancing compound is about 0.2 to 2 mg/ml.
- 34. (original) The composition according to claim 28, wherein the agent modulates a biological process in a cell when the agent is present in the cell.
- 35. (original) The composition according to claim 34, wherein the biological process is selected from the group consisting of cell growth, differentiation, proliferation, a metabolic or biosynthetic pathway, gene expression, a disease-associated process, and an immune response.
- 36. (original) The composition according to claim 28, wherein the agent comprises a polynucleotide.
- 37. (original) The composition according to claim 36, wherein the polynucleotide is selected from the group consisting of an antisense nucleic acid, a triplex-forming nucleic acid, and a nucleic acid that comprises a gene which encodes a polypeptide.
- 38. (original) The composition according to claim 37, wherein the gene is a tumor suppressor gene.
- 39. (original) The composition according to claim 37, wherein the tumor suppressor gene is selected from the group consisting of a retinoblastoma gene and a p53 gene.
- 40. (original) The composition according to claim 28, wherein the composition further comprises a polymeric matrix.
- 41. (original) The composition according to claim 28, wherein the composition further comprises a mucoadhesive.

42. (previously presented) A delivery enhancing compound having a

Formula I:

$$X_1$$
— C — N — $(CH_2)_m$ — N — $(CH_2)_n$ — N — R
 C = O
 X_2

wherein:

m and n are the same or different and each is an integer from 2-8; R [[is]] forms a cationic group with the nitrogen to which it is bound, or

 X_1 is selected from the group consisting of:

 X_2 , and X_3 are each independently selected from the group consisting of a saccharide group,

wherein at least one of X2 and X3 is a saccharide group when R is

- 43. (previously presented) The compound of claim 42, wherein R forms a cationic group selected from the group consisting of NMe₃⁺ and NH₃⁺.
- 44. (previously presented) The compound of claim 42, wherein the saccharide group has between one to eight monosaccharide groups.
- 45. (original) The compound of claim 44, wherein the saccharide group is selected from the group consisting of pentose monosaccharide groups, hexose monosaccharide groups, pentose-pentose disaccharide groups, hexose-hexose disaccharide groups, pentose-hexose disaccharide groups, and hexose-pentose disaccharide groups.
- 46. (original) The compound of claim 42, wherein the saccharide group comprises between three and about eight monosaccharide residues.
- 47. (original) The compound of claim 46, wherein the saccharide group is a trisaccharide.
- 48. (original) The compound of claim 42, wherein at least one of X_2 and X_3 is a saccharide group.
- 49. (original) The compound of claim 42, wherein m and n are each independently 2 or 3.

both

50. (original) The compound of claim 42, wherein both X_1 and X_2 are

and X₃ is a saccharide group.

- 51. (original) The compound of claim 42, wherein the saccharide group is a hexose-hexose disaccharide group.
- 52. (original) The compound of claim 42, wherein m and n are each 3, X_1 and X_2 are both

and X₃ is a hexose monosaccharide group.

53. (original) The compound of claim 42, wherein m and n are each 3, X₁ and X₃ are both

and X_2 is a hexose monosaccharide group.

54. (original) The compound of claim 42, wherein m and n are each 3, X_1 and X_2 are both

and X₃ is a hexose-hexose disaccharide group.

55. (original) The compound of claim 42, wherein m and n are each 3, X_1 and X_3 are both

X₂ is a hexose-hexose disaccharide group.

56. (currently amended) The compound composition according to claim 42 28, wherein the compound has a Formula III:

57. (currently amended) The compound composition according to claim 42 28, wherein the compound has a Formula IV:

58. (currently amended) The compound composition according to claim 42 28, wherein the compound has a Formula V:

59-81. (Canceled)

- 82. (previously presented) The composition according to claim 28, wherein the agent is a gene encoding interferon.
- 83. (previously presented) The composition according to claim 82, wherein the interferon is a member of the group selected from α -interferon, β -interferon, δ -interferon, and γ interferon.
- 84. (previously presented) The composition according to claim 83, wherein the interferon is α -interferon.

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- 85. (currently amended) The composition according to claim 83 28, wherein the gene is incorporated into a vector.
- 86. (currently amended) The composition according to claim 83 28, wherein the vector is a recombinant viral vector.
- 87. (currently amended) The composition according to claim 83 86, wherein the recombinant viral vector is selected from the group consisting of a herpes viral vector, retroviral vector, vaccinia viral vector and an adenoviral vector
- 88. (previously presented) The composition according to claim 87, wherein the recombinant viral vector is an adenoviral vector.